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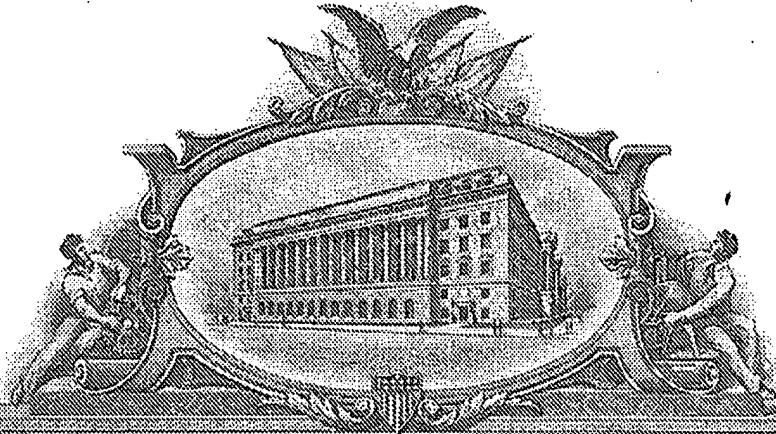
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PROVISIONAL APPLICATION FOR PATENT COVER SHEET

This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53(c).

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07/01/03

INVENTOR(S)					
Given Name (first and middle (if any))		Family Name or Surname		Residence (City and either State or Foreign Country)	
Jun		Hu		Fairlawn, OH	
<input type="checkbox"/> Additional inventors are being named on the _____ separately numbered sheets attached hereto					
TITLE OF THE INVENTION (500 characters max)					
METHYL-2,4,9-TRITHIAADAMANTANE-7-CARBOXYLATE					
Direct all correspondence to: CORRESPONDENCE ADDRESS					
<input type="checkbox"/> Customer Number		Type Customer Number here		Place Customer Number Bar Code Label here	
OR					
<input checked="" type="checkbox"/> Firm or Individual Name		Roetzel & Andress			
Address		222 South Main St.			
Address					
City	Akron	State	Ohio	ZIP	44308
Country	US	Telephone	330-849-6777	Fax	330-376-4577
ENCLOSED APPLICATION PARTS (check all that apply)					
<input checked="" type="checkbox"/> Specification	Number of Pages	9	<input type="checkbox"/> CD(s), Number		
<input checked="" type="checkbox"/> Drawing(s)	Number of Sheets	4	<input type="checkbox"/> Other (specify)		
<input type="checkbox"/> Application Data Sheet. See 37 CFR 1.76					
METHOD OF PAYMENT OF FILING FEES FOR THIS PROVISIONAL APPLICATION FOR PATENT					
<input checked="" type="checkbox"/>	Applicant claims small entity status. See 37 CFR 1.27.			FILING FEE AMOUNT (\$)	
<input type="checkbox"/>	A check or money order is enclosed to cover the filing fees				
<input checked="" type="checkbox"/>	The Commissioner is hereby authorized to charge filing fees or credit any overpayment to Deposit Account Number: 50-0959(089498-0452)			\$80.00	
<input type="checkbox"/>	Payment by credit card. Form PTO-2038 is attached.				
The invention was made by an agency of the United States Government or under a contract with an agency of the United States Government.					
<input checked="" type="checkbox"/> No.					
<input type="checkbox"/> Yes, the name of the U.S. Government agency and the Government contract number are: _____					

Respectfully submitted,

SIGNATURE

TYPED or PRINTED NAME Daniel J. Schlue

TELEPHONE 330-849-6777

Date

07/01/2003

REGISTRATION NO.

(if appropriate)

Docket Number:

52,194

089498-0452

USE ONLY FOR FILING A PROVISIONAL APPLICATION FOR PATENT

This collection of information is required by 37 CFR 1.51. The information is used by the public to file (and by the PTO to process) a provisional application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 8 hours to complete, including gathering, preparing, and submitting the complete provisional application to the PTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, Washington, D.C. 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Box Provisional Application, Assistant Commissioner for Patents, Washington, D.C. 20231.

METHYL 2,4,9-TRITHIAADAMANTANE-7-CARBOXYLATE**FIELD OF THE INVENTION**

5 This invention relates to methyl 2,4,9-trithiaadamantane-7-carboxylate and a method for its manufacture.

BACKGROUND OF THE INVENTION

10 Growing an α -helical peptide from a substrate coated with a metallic film is known. An exemplary metallic film is made of gold, and the α -helical peptide is attached to the metallic film by a surface anchor. The anchor functions by chemically bonding with both the peptide and the film surface- thereby securing one to the other.

15 Prior-art anchors ordinarily bond to a metallic-film surface via a single sulfur-metal bond. In other words, prior-art anchors ordinarily have a single sulfur functionality that is reactive with a metallic-film surface. The deficiency of such anchors is that their resultant surface orientation is often uncontrollable and unpredictable. For example, their positioning is either substantially vertical or substantially parallel to the metallic-film surface. And that, in turn, directly impacts the spatial orientation of an attached α -helical peptide. So if an anchor's spatial orientation is substantially parallel to a film surface, the attached α -helical peptide is also positioned substantially parallel to the film surface. Uncontrollable surface
20 orientation is problematic because an α -helical peptide functions best as an optical switch when it is substantially vertical. The anchor's spatial orientation is therefore key.

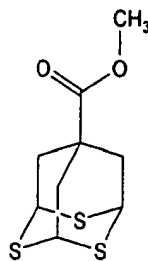
In addressing the need for sulfur-based surface anchors having predictable surface orientation, the prior art teaches using a trithiaadamantane anchor to grow an α -helical peptide. Trithiaadamantane has a three-dimensional tripodal structure that provides more predictable
25 surface orientation compared to single-sulfur anchors. Trithiaadamantane anchors form three sulfur-metal bonds with a metal surface, and that causes the compound to assemble in an upright vertical position. The anchor's upright surface orientation causes the α -helical peptide to likewise have a vertical spatial orientation, and that is preferred.

Prior-art trithiaadamantane anchors have previously been manufactured only by using the specific chemical intermediate- ethyl 2,4,9-trithiaadamantane-7-carboxylate. The prior art fails to provide useful alternate intermediates. A significant drawback to any method that employs ethyl-2,4,9-trithiaadamantane-7-carboxylate is that ethyl-2,4,9-trithiaadamantane-7-carboxylate can only be produced at the relatively low yield of about 10-25%.

So the art needs an alternate intermediate compound, one other than ethyl 2,4,9-trithiaadamantane-7-carboxylate, that can be used to produce trithiaadamantane anchors for growing α -helical peptides. There is also a need for an alternate intermediate compound that can be used in making trithiaadamantane anchors, wherein the alternate intermediate compound can be produced at a yield greater than the prior art's 10-25% yield for ethyl-2,4,9-trithiaadamantane-7-carboxylate. There is an additional need for methods directed to making trithiaadamantane anchors.

SUMMARY OF THE INVENTION

This invention provides a compound having the formula:



This invention further provides a method for synthesizing methyl 2,4,9-trithiaadamantane-7-carboxylate comprising the step of reacting oxidized methyl triallyl acetate with a sulphuring agent and a Lewis acid to produce methyl 2,4,9-trithiaadamantane-7-carboxylate.

Some advantages of this invention are that methyl 2,4,9-trithiaadamantane-7-carboxylate can be employed as a chemical intermediate in manufacturing compounds having trithiaadamantane anchors. Advantageously, methyl 2,4,9-trithiaadamantane-7-carboxylate is

a drop-in replacement for ethyl-2,4,9-trithiaadamantane-7-carboxylate in methods for making compounds having trithiaadamantane surface anchors. A noted advantage over the prior-art intermediate, ethyl-2,4,9-trithiaadamantane-7-carboxylate, is that methyl 2,4,9-trithiaadamantane-7-carboxylate can be synthesized in a 35-40% yield, which is a clear improvement over the 10-25% yield of prior-art methods for producing ethyl-2,4,9-trithiaadamantane-7-carboxylate.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a ^1H NMR spectrum of methyl 2,4,9-trithiaadamantane-7-acetate produced by the subject method.

Fig. 2 is a ^1H NMR spectrum of 2,4,9-trithiaadamantane-7-carboxylic acid produced by the subject method.

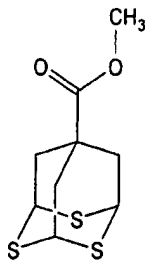
Fig. 3 is a ^{13}C NMR spectrum of methyl 2,4,9-trithiaadamantane-7-carboxylate produced by the subject method.

Fig. 4 is an IR spectrum of methyl 2,4,9-trithiaadamantane-7-carboxylate produced by the subject method.

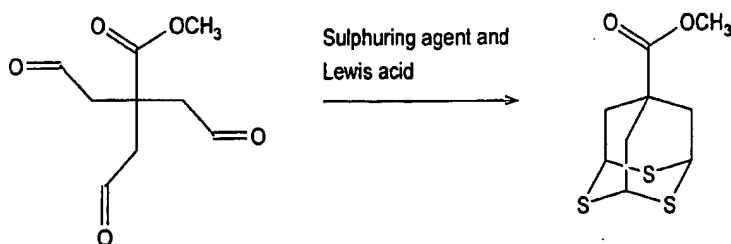
DETAILED DESCRIPTION OF THE INVENTION

This invention is generally directed to methyl 2,4,9-trithiaadamantane-7-carboxylate and a method for its manufacture.

Methyl 2,4,9-trithiaadamantane-7-carboxylate is represented by the formula:



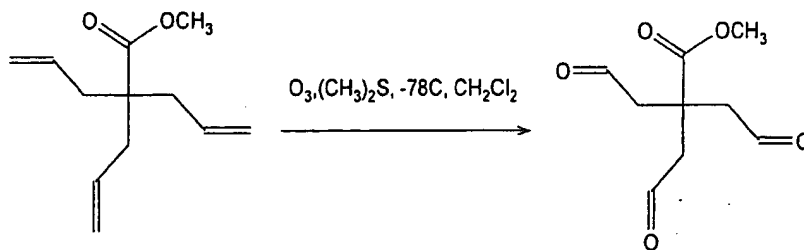
Methyl 2,4,9-trithiaadamantane-7-carboxylate is synthesized by reacting oxidized methyl triallyl acetate with a sulphuring agent and a Lewis acid. A general understanding of the method can be gained from the following simplified reaction scheme:



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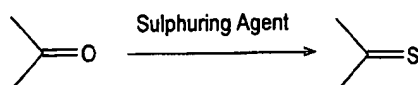
A nonlimiting example of this method adds Lawesson's reagent and $\text{BF}_3 \cdot \text{Et}_2\text{O}$ to oxidized methyl triallyl acetate. The reaction then proceeds under reflux for approximately 100 hours.

Oxidized methyl triallyl acetate is preferably synthesized, and that is accomplished by the oxidation of methyl triallyl acetate. Any known method of oxidation can be employed. But ozonolysis is preferred. Ozonolysis of methyl triallyl acetate can be performed by preparing a solution of methyl triallyl acetate in freshly-distilled methylenechloride followed by stirring and cooling the solution to -78°C in a dry-ice acetone bath. Ozone is subsequently bubbled through the solution until a light-blue color persists. A simplified reaction scheme is provided:



15

Any sulphuring agent can be employed in synthesizing methyl 2,4,9-trithiaadamantane-7-carboxylate. A sulfuring agent converts ketonic groups into thioketonic groups as represented by the reaction scheme:



5 Preferred sulphuring agents include 2,4-bis(4-methoxyphenyl)-1,3,2,4-dithiadiphosphetane 2,4-disulfide (Lawesson's reagent), phosphorous pentasulfide hexamethyldisiloxane (PPHD), or combinations thereof.

10 The relative mole ratio of sulphuring agent to oxidized methyl triallyl acetate can generally range from about 6:1 to about 1:1. Preferably, the mole ratio ranges from about 3:1 to about 2:1.

Any Lewis acid can be employed in practicing the present invention. Preferred Lewis acids include the complex of boron trifluoride and ethyl ether ($\text{BF}_3 \cdot \text{Et}_2\text{O}$), the complex of boron trichloride and ethyl ether ($\text{BCl}_3 \cdot \text{Et}_2\text{O}$), or combinations thereof.

15 The mole ratio of Lewis acid to oxidized methyl triallyl acetate preferably ranges from about 4:1 to about 1:1. More preferably, the mole ratio ranges from about 3:1 to about 2:1. Generally, the concentration of Lewis acid in the reaction medium ranges from about 0.5 to about 1.0 molar (*M*).

20 Any noncoordinated or weakly coordinated reaction solvent can be employed in synthesizing methyl 2,4,9-trithiaadamantane-7-carboxylate. Nonlimiting examples of preferred solvents are methylene chloride (CH_2Cl_2), carbon tetrachloride (CCl_4), benzene, or combinations thereof. Effective amounts of non-coordinated or weakly coordinated solvent can be determined by a person of ordinary skill in the art without undue experimentation.

25 Synthesizing methyl 2, 4, 9-trithiaadamantane-7-carboxylate preferably occurs under reflux, and persons of ordinary skill in the art can determine the required temperature without undue experimentation.

EXAMPLES

In order to demonstrate practice of the present invention, the following examples have been prepared and tested. The examples should not, however, be viewed as limiting the scope of the invention. The claims will serve to define the invention.

5

Methyl 2,4,9-trithiaadamantane-7-carboxylate

A solution of 5 grams of methyl triallyl acetate in 100 milliliters freshly distilled methylenechloride was stirred and cooled to -78°C in a dry-ice acetone bath. Ozone was bubbled through the cooled mixture until a light-blue color persisted. The ozone line was then disconnected and the excess ozone was removed by argon flow for 10 minutes. 6.0 grams dimethyl sulfide was added to the reaction mixture at -78°C. The mixture was slowly warmed up to the ambient temperature. The mixture was then concentrated via rotary evaporation. To this mixture, 100 milliliters of methylenechloride, 28.0 grams of Lawesson's reagent, and 100 milliliters of neat $\text{BF}_3 \cdot \text{Et}_2\text{O}$ were added respectively. The mixture was then refluxed for 100 hours. Additional methylenechloride was added (100 mL). The mixture was washed using conventional techniques with 0.2 M potassium carbonate three times. The organic layer was dried over magnesium sulfate and evaporated. The resulting mixture was purified by column chromatography on silica gel using 30% methylenechloride in hexane to give pure methyl 2,4,9-trithiaadamantane-7-carboxylate (35-40% yield; melt point 149°C-151°C).

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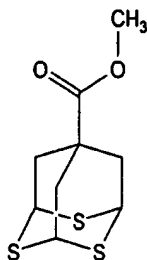
It should be evident that the present invention substantially improves the art by providing methyl 2,4,9-trithiaadamantane-7-carboxylate and a method for making it. While, in accordance with the patent statutes, only the preferred embodiments of the present invention have been described in detail hereinabove, the present invention is not to be limited thereby. Rather, the scope of the invention shall include all modifications and variations that fall within the scope of the attached claims.

25

CLAIMS

What is claimed is:

1. A compound having the formula:



- 5 2. A method for synthesizing methyl 2,4,9-trithiaadamantane-7-carboxylate comprising of the step of:
reacting oxidized methyl triallyl acetate with a sulphuring agent and a Lewis acid to produce methyl 2,4,9-trithiaadamantane-7-carboxylate.
- 10 3. The method of claim 2, wherein the sulphuring agent is 2,4-bis(4-methoxyphenyl)-1,3,2,4-dithiadiphosphetane 2,4-disulfide, phosphorus pentasulfide hexamethyldisiloxane, or a combination thereof.
- 15 4. The method of claim 2, wherein the sulphuring agent is reacted with oxidized methyl triallyl acetate in a relative mole ratio ranging from about 6:1 to about 1:1.
5. The method of claim 2, wherein the sulphuring agent is reacted with oxidized methyl triallyl acetate in a relative mole ratio ranging from about 3:1 to about 2:1
- 20 6. The method of claim 2, wherein the Lewis acid is $\text{BF}_3 \cdot \text{Et}_2\text{O}$, $\text{BCl}_3 \cdot \text{Et}_2\text{O}$, or a combination thereof.

7. The method of claim 2, wherein the Lewis acid is reacted with the oxidized methyl triallyl acetate in a relative mole ratio ranging from about 4:1 to about 1:1.
8. The method of claim 2, wherein the Lewis acid is reacted with the oxidized methyl triallyl acetate in a relative mole ratio ranging from about 3:1 to about 2:1.
9. The method of claim 2, wherein the step of reacting oxidized methyl triallyl acetate with a sulphuring agent and a Lewis acid to produce methyl 2,4,9-trithiaadamantane-7-carboxylate occurs in a noncoordinated or weakly coordinated solvent.
10. The method of claim 9 wherein the noncoordinated or weakly coordinated solvent is methylene chloride, carbon tetrachloride, benzene, or a combination thereof.

Docket No: 089498.0452

ABSTRACT OF THE DISCLOSURE

Methyl 2,4,9-trithiaadamantane-7-carboxylate and a method for its manufacture is disclosed. The method reacts oxidized methyl triallyl acetate with a Lewis acid and a sulphuring agent.

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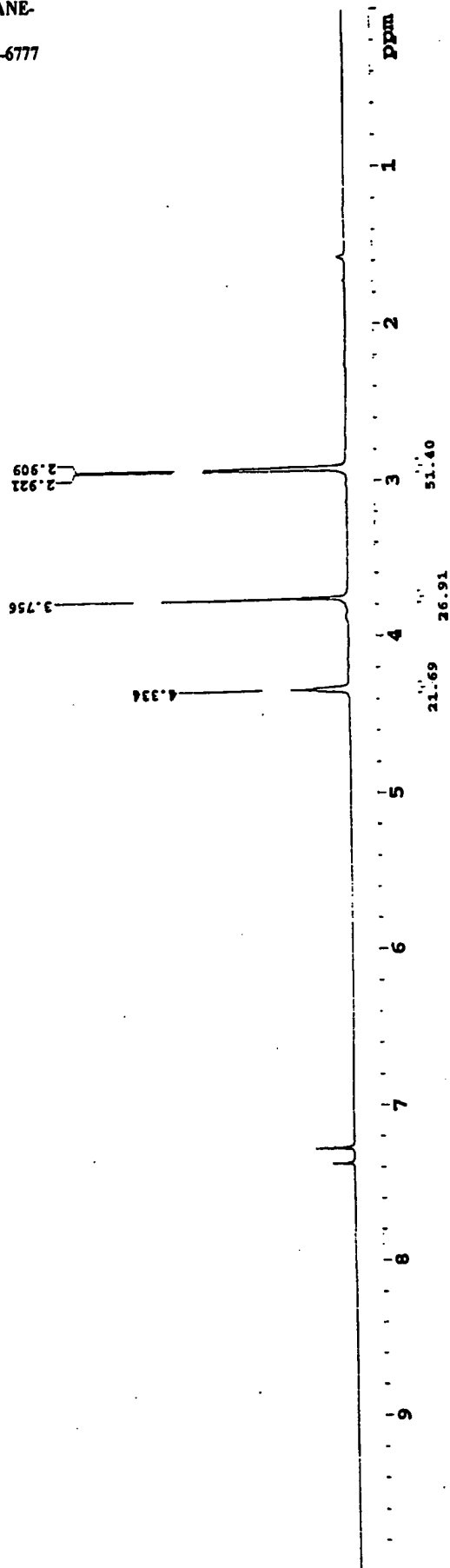


FIG. 1

Jun Hu 089498-0452
METHYL 2,4,9-TRITHIAADAMANTANE-
7-CARBOXYLATE
Daniel J. Schlue 330-849-6777
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Page 2 of 4

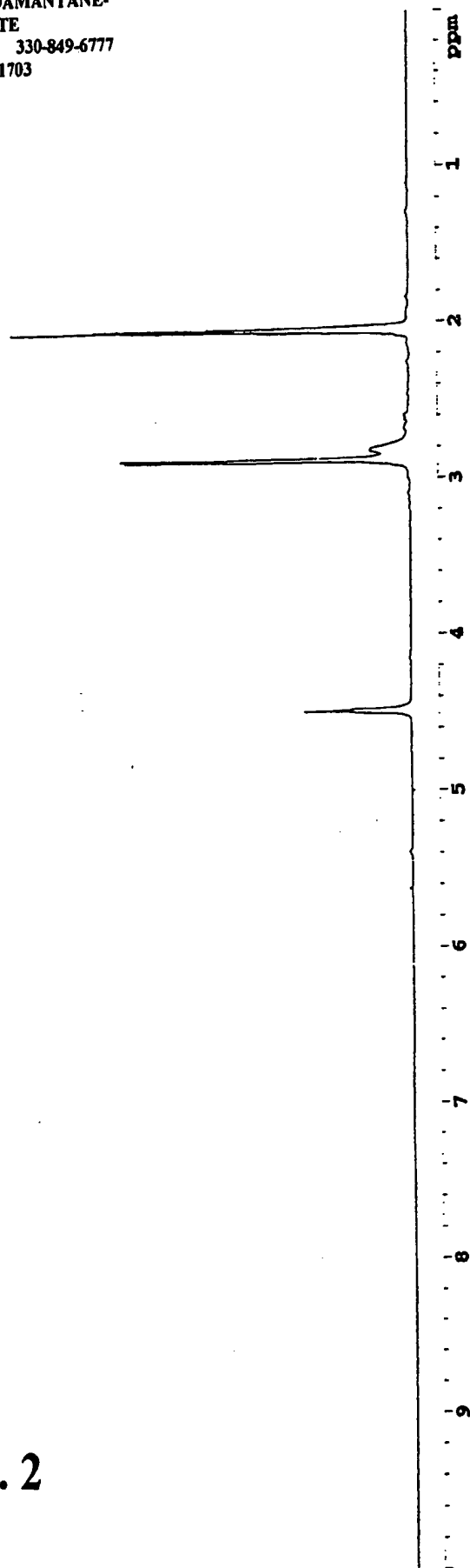


FIG. 2

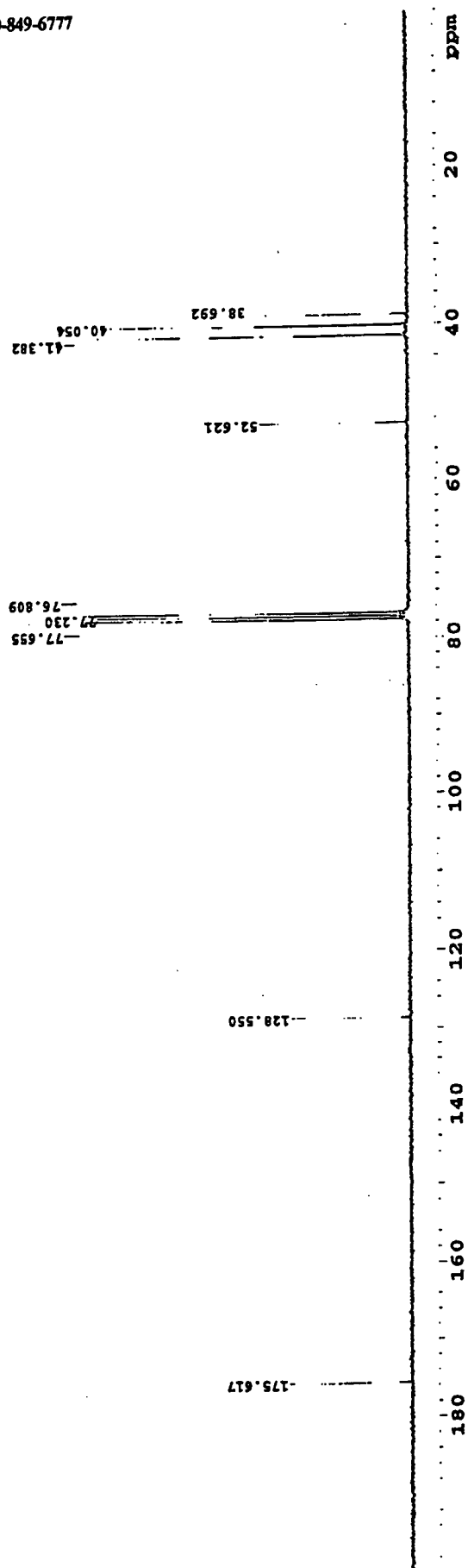


FIG. 3

Jun Hu 089498-0452
METHYL 2,4,9-TRITHIAADAMANTANE-

7-CARBOXYLATE
Daniel J. Schlue 330-849-6777
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Page 4 of 4

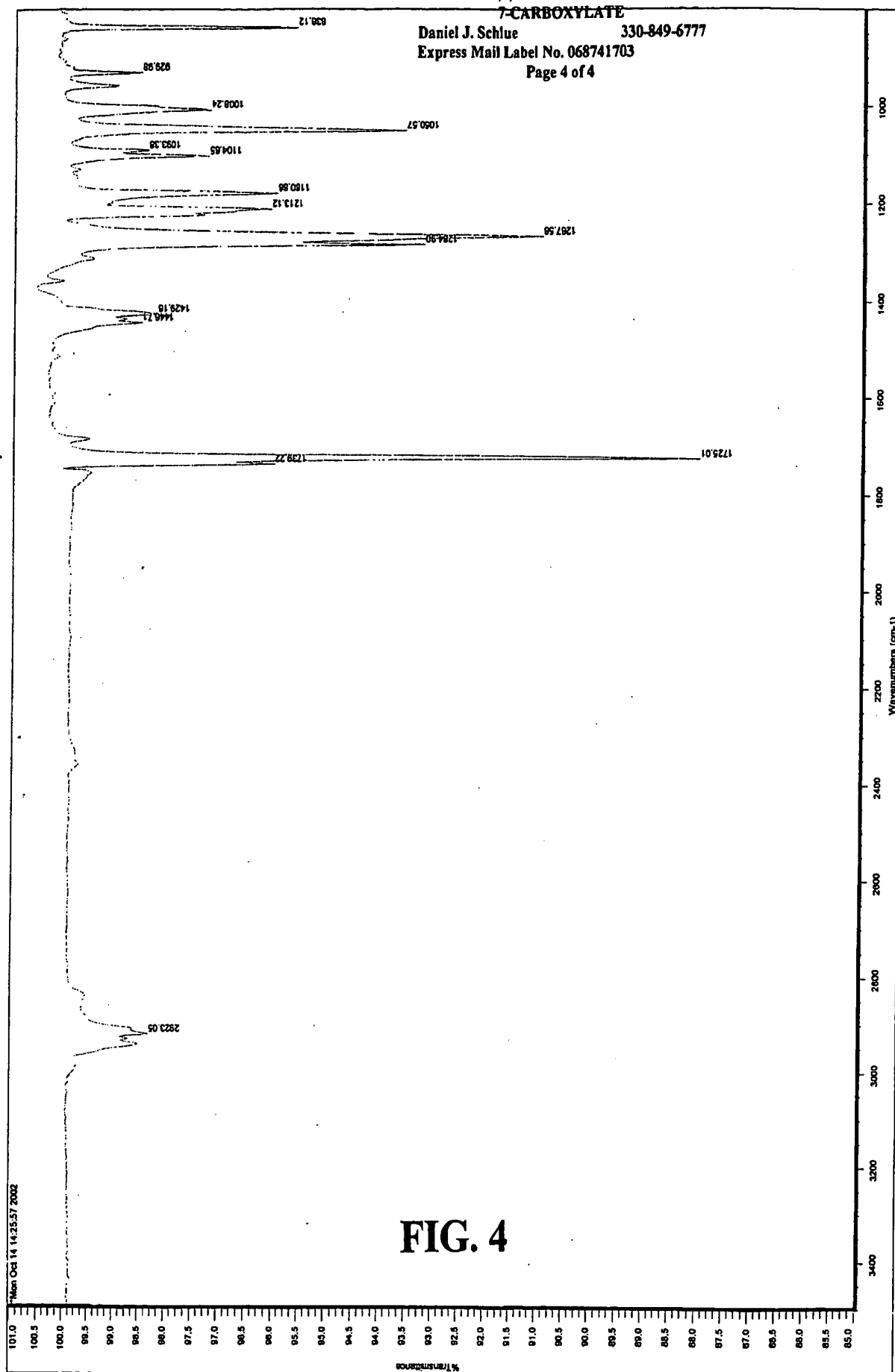


FIG. 4